- **68-7-14. Prescription labels.** (a) The label of each drug or device shall be typed or machine-printed and shall include the following information:
- (1) The name, address, and telephone number of the pharmacy dispensing the prescription;
- (2) the name of the prescriber. The label shall include the name of the practitioner and, if involved, the name of either the physician's assistant (PA) or the advanced registered nurse practitioner (ARNP);
  - (3) the full name of the patient;
- (4) the identification number assigned to the prescription by the dispensing pharmacy;
  - (5) the date the prescription was filled or refilled;
  - (6) adequate directions for use of the drug or device;
  - (7) the beyond-use date of the drug or device dispensed;
  - (8) the brand name or corresponding generic name of the drug or device;
- (9) the name of the manufacturer or distributor of the drug or device, or an easily identified abbreviation of the manufacturer's or distributor's name;
  - (10) the strength of the drug;
  - (11) the contents in terms of weight, measure, or numerical count; and
  - (12) necessary auxiliary labels and storage instructions, if needed.
- (b) A pharmacy shall be permitted to label or relabel only those drugs or devices originally dispensed from the providing pharmacy. (Authorized by

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